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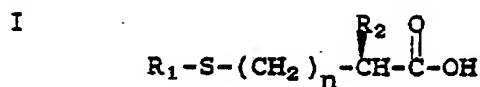
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(54) Enzymatic resolution process.

(57) S-Enantiomers of the formula



result from the reaction of a racemic form of compound I with an alcohol in the presence of an enzyme capable of selectively esterifying the R-enantiomers of the racemic compound..

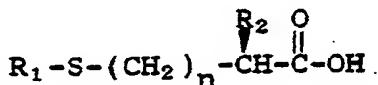
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ENZYMATIC RESOLUTION PROCESS

In accordance with the present invention a novel process for preparing S-enantiomers having the formula

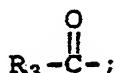
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I



where R_1 is hydrogen or

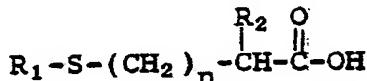
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15 R_2 and R_3 are each independently selected from alkyl; cycloalkyl aralkyl or aryl; and n is 1 or 2, is provided. The present process comprises reacting a racemic mixture of a compound having the formula

20

I'



with a compound of the formula

VIII R_4-OH ,
25 wherein R_4 is alkyl, cycloalkyl, aryl or aralkyl, in the presence of an enzyme or microorganism capable of catalyzing the selective esterification of compounds of formula I' to provide a solution containing R-enantiomeric esters and S-enantiomeric unreacted acids of formula I; and, recovering the resulting non-esterified S-enantiomers of formula I.

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Detailed Description of Preferred Embodiments

The following definitions apply throughout this application.

35 The term "alkyl" as used herein refers to straight or branched chain carbon groups of 1 to 25 carbon atoms, preferably 1 to 6 carbon atoms.

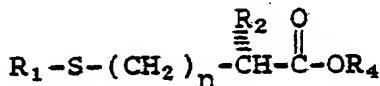
The term "cycloalkyl" as used herein refers to groups containing 5 to 7 carbon atoms.

40 The term "aryl" as used herein refers to monocyclic or bicyclic aromatic groups containing from 6 to 10 carbon atoms in the ring portion such as phenyl, naphthyl, and substituted phenyl or naphthyl containing substituents such as nitro, halogen, methyl or alkoxy groups on the aromatic ring.

More specifically, it has been found that alcohols of formula VIII, in the presence of catalyzing lipases or esterases (or microorganisms producing same), are capable of catalyzing the stereoselective esterification of racemic compounds of formula I'. This process produces esters in the R-enantiomeric form having the formula

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IX

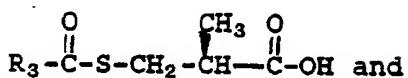


50 and the resulting "by-product" is in fact, a high yield of optically pure unreacted S-enantiomers of formula I.

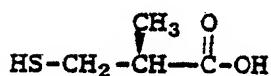
The enzymatic resolution process of the present invention has the advantage that it can provide the desired S-enantiomers of formula I with high optical purity, and at good yields. Additionally, because the present process uses a racemic carboxylic acid of formula I' as a starting material, instead of the carboxylic acid esters employed in prior art enzymatic processes, there is considerably less expense involved. These

and other features make the process of the present invention very attractive for use in preparing optically active compounds of formula I, such as the S-enantiomers of the formula

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II

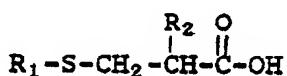
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IIa

useful in the preparation of captopril and zofenopril.

Methods for obtaining the racemic starting material of the formula

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I

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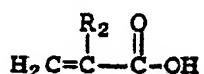
are known.

For example, a compound of the formula

VII $\text{R}_1\text{-SH}$

can be coupled to a compound of the formula

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VIII

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in the presence or absence of a suitable solvent, such as hexane, heptane or isopropanol, under the usual conditions for conducting such an addition reaction.

Compounds of formula

VIII $\text{R}_4\text{-OH}$

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for use in the present process can be any alcohol suitable for esterifying compounds of formula I. For example, in a preferred embodiment for providing resolved S-enantiomers of formula II and IIa, methanol, benzyl alcohol and 1-octanol each provided optical purities in excess of 95% at reaction yields of between about 30% and 40%.

40

The present process can be carried out in an organic solvent. Typical solvents suitable for use in the present process include, but are not limited to, 1,1,2-trichloro-1,2,2-trifluoroethane, toluene, cyclohexane, benzene, hexane, heptane, isoctane and octane.

45

The enzyme or microorganism used in the present process can be any enzyme or microorganism having the ability to catalyze the stereoselective esterification of acids of formula I. Various enzymes, such as esterases and lipases, regardless of origin or purity, are suitable for use in the present invention. The enzyme can be in the form of a mixture of animal and plant enzyme, cells of microorganisms, crushed cells or extracts of cells.

50

Typical genera of microorganism suitable as sources of catalyzing enzymes include *Mucor*, *Escherichia*, *Staphylococcus*, *Agrobacterium*, *Rhizopus*, *Aspergillus*, *Nocardia*, *Streptomyces*, *Trichoderma*, *Candida*, *Rhodotorula*, *Torylopsis*, *Bacillus*, *Alcaligenes*, *Pseudomonas*, *Brevibacterium*, *Enterobacter*, *Chromobacterium*, *Arthrobacter*, *Microbacterium*, *Mycobacterium*, *Saccharomyces*, *Penicillium*, *Botrytis*, *Chaetomium*, *Ophiobolus*, *Cladosporium* and the like.

55

Commercially available enzymes suitable for use in the present invention include lipases, such as Amano P (*Pseudomonas fluorescens*) which is preferred, Amano AY-30 (*Candida cylindracea*), Amano N (*Rhizopus niveus*), Amano R (*Penicillium sp.*), Amano FAP (*Rhizopus oryzae*), Amano AP-12 (*Aspergillus niger*), Amano MAP (*Mucor meihai*), Amano GC-4 (*Geotrichum candidum*), Sigma L-0382 (porcine pancreas), Sigma L-3001 (Wheat germ), Sigma L-1754 (*Candida cylindracea*), Sigma L-0763 (*Chromobacterium viscosum*) and Amano K-30 (*Aspergillus niger*). Additionally, enzymes derived from animal tissue include esterase from pig liver, α -chymotrypsin and pancreatin from pancreas.

Specific microorganisms suitable for use in the present process include *Pseudomonas fluorescens*,

Pseudomonas putida, Pseudomonas ovalis, Escherichia coli, Staphylococcus aureus, Alcaligenes faecalis, Streptomyces griseus, Streptomyces clavuligervs, Nocardia ethropolis, Nocardia asteroides, Mycobacterium phlei, Agrobacterium radiobacter, Aspergillus niger, Rhizopus oryzae and the like.

In the preferred process of the present invention, the alcohol of formula VIII and racemic starting material of formula I are added to the desired organic solvent. The enzyme (or microorganism containing same) is added thereto. Typically, the enzyme is adsorbed onto a suitable carrier, e.g. diatomaceous earth (porous Celite Hyflo Supercel), microporous polypropylene (Enka Accurel® polypropylene powder), or a nonionic polymeric adsorbent such as Amberlite® XAD-2 (polystyrene) or XAD-7 (polyacrylate) from Rohm and Haas Co. This serves the purpose of immobilizing the enzyme which has the effects of controlling the enzyme particle size and preventing aggregation of the enzyme particles when used in an organic solvent. This can be accomplished, for example, by precipitating an aqueous solution of the enzyme with cold acetone in the presence of the Celite Hyflo Supercel followed by vacuum drying, or in case of nonionic polymeric adsorbent, incubating enzyme solutions with adsorbent on a shaker for desired time, removing excess solution and drying enzyme-adsorbent resins under vacuum. The reaction solution typically contains between about 5 and 250 mg of racemic starting material per ml of solvent. The enzyme added to the reaction solution may be present in concentrations ranging from about 5 to about 200 mg of enzyme per ml of solvent. While it is desirable to use the least amount of enzyme possible, the amount of enzyme required will vary depending upon the specific activity of the enzyme used.

When the reaction is conducted in an organic solvent, small amounts of water may be added to the reaction mixture. The water added to the reaction mixture may be present in concentrations ranging from about 0.2 to about 100 mg of water per ml of solvent, or solvent saturated with water, and preferably is present in an amount of about 0.5-5 mg/ml. Incubation of the reaction solution can be at a temperature between about 4 and about 60 °C and is preferably carried out at about 30-50 °C. The reaction time can be appropriately varied depending upon the amount of enzyme used and its specific activity. Typical reaction times at 28 °C for optical purities of 90 percent and above are at least about 2 hours and can range up to about 96 hours for greater conversions and higher optical purities, e.g. optical purities exceeding 95 percent. Reaction times can be reduced significantly by increasing the reaction temperature and/or increasing the amount of enzyme added to the reaction solution. S-enantiomers of formula I can be isolated from the reaction mixture and purified by known methodologies such as extraction, distillation, crystallization, column chromatography, and the like.

As will be apparent to those skilled in the art, the process of the present invention can be carried out using microbial cells containing an enzyme having the ability to catalyze the stereo-selective esterification of compounds of formula I. When using a microorganism to perform the resolution, the present process is conveniently carried out by adding the cells and the racemic starting material to the desired solvent. Cells may be used in the form of intact cells, dried cells such as lyophilized, spray-dried or heat-dried cells, immobilized cells, or cells treated with organic solvents such as acetone or toluene. Cells may also be used in the form of treated cell material such as ruptured cells or cell extract.

Using the methodology of U.S. 4,105,776, the resolved acid of formula I or its chemical equivalent is used to acylate L-proline having the formula

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VIII



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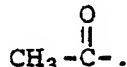
forming captopril, i.e. the compound of formula III, in the case where R₁ in compound I is hydrogen. Preferably R₁ in compound I is an

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group, such as

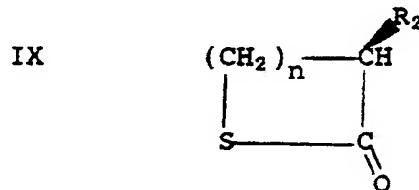
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In this case, the thioester formed by the coupling of compound I with compound VIII can be deacylated by

conventional means, such as by ammonolysis (e.g., by treatment with alcoholic ammonia or concentrated ammonium hydroxide) or by alkaline hydrolysis (e.g., by treatment with aqueous metal hydroxide). Alternatively, again employing methodology from U. S. 4,105,776, the resolved acid of formula I can be used by removing the acetyl group (by conventional methods) and thereafter dehydrating the so-treated acid to form a thiolactone of the formula

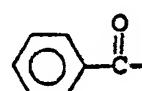
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15 The thiolactone IX can thereafter be used to acylate the L-proline of formula VIII to obtain the desired product.

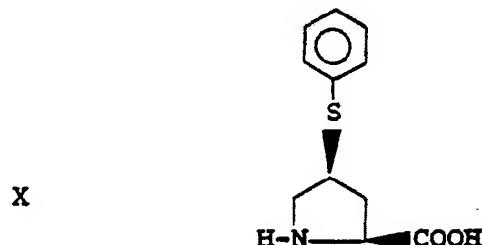
Similarly, to provide zofenopril, i.e., the compound of formula IV, the resolved acid of formula I or its chemical equivalent where R₁ is

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25 is used to acylate a compound of the formula

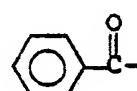
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as described in U. S. 4,316,906. Alternatively, using the methodology of U. S. 4,316,906, the compound of formula I where R₁ is

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45 can be coupled with a compound of the formula

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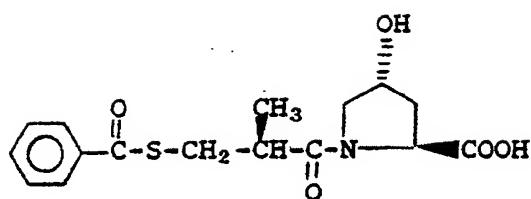


or esters or protected forms thereof, to provide a compound of the formula

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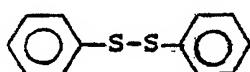
XII



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which can thereafter be treated with a compound of the formula

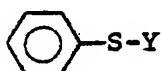
XIII



15

or

XIV



20

where Y is an activating group such as succinimide or phthalamido or a halide such as Cl or Br to provide the product of formula IV.

The acylation of compounds VIII, X or XI with the resolved acid of formula I can be effected in the presence of a coupling agent like dicyclohexycarbodiimide or the like, or the acid can be activated by formation of its mixed anhydride, symmetrical anhydride, acid chloride, acid ester or use of Woodward reagent K, N-ethoxy-carbonyl-2-ethoxy-1,2-dihydroquinoline or the like. For a review of the methods for acylation, see Methoden der Organischen Chemie (Houben-Weyl), Vol. XV, part II, page 1 et seq. (1974).

The present invention will now be described by the following examples, however, it should be understood that the invention is not meant to be limited by the details therein.

Example 1

35 Racemic 3-benzoylthio-2-methylpropanoic acid (SQ 32,979; 6.726 g; 0.03 mol; 22.42 mg/ml) and methanol (3.845 g, 0.12 mol) were dissolved in and diluted with toluene to 300 ml. Crude Amano lipase P-30 (12 g; 40 mg/ml; *Pseudomonas fluorescens*) and deionized water (300 µl) were added and the reaction mixture was incubated at 28 °C on a shaker at 280 RPM. The esterification was monitored by capillary gas chromatography (GC) with a flame ionization detector (FID) to determine reaction yields. At 198 hours 2.475 g of the remaining unreacted substrate, that is,

40 S(-)-3-benzoylthio-2-methylpropanoic acid (SQ 25,680), had an optical purity of 97.1% with a reaction yield of 37% (results shown in Table 1). In this and all following examples, the enantiomeric composition of the unreacted 3-benzoylthio-2-methylpropanoic acid fraction was determined by capillary gas chromatography following derivatization with thionyl chloride and esterification of the resulting acid chloride with (S)-(+) -2-

45 octanol to form diastereomeric esters which can be separated by capillary GC.

Examples 2-4

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Following the procedure of Example 1, additional resolutions were carried out with methanol, benzyl alcohol and 1-octanol. The results are summarized in Table 2.

55

Table 1

Enzymatic Resolution of SQ 25,680						
	Reaction Time (hr)	Conversion (%)	Reaction Yield (%)	Optical Purity (%)	e.e. (%)	Enantiomeric Ratio E
5	0	0	100	50.0	0.0	
10	136	57	43	92.3	84.6	12
15	159	58	42	95.0	90.0	15
20	166	59	41	95.5	91.0	14
25	198	63	37	97.1	94.2	12
Reaction system: 0.1 M SQ 32,979 (22.42 mg/ml), 0.4 M methanol in 300-ml toluene; 300 uL (0.1 %v/v) deionized H ₂ O added to reaction mixture.						
Enzyme: 12.0 g (40 mg/ml) Amano lipase P-30 (from <i>Pseudomonas fluorescens</i>)						
Conditions: 28 °C, 280 rpm.						

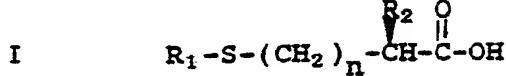
Table 2

Ex. No.	Alcohol	Reaction Time (hr)	Conversion (%)	Reaction Yield (%)	Optical Purity (%)	e.e. (%)	Enantiomeric Ratio E	
25	2	0	0.0	100.0	50.0	0.0		
		16	5.8	94.2				
		40	33.5	66.5	63.8	27.6	4.5	
		112	50.3	49.7	91.7	83.4	27.0	
		136	55.1	44.9	95.4	90.8	20.7	
		162	63.0	37.0	97.3	94.6	11.9	
		280	76.5	23.5	97.7	95.4	5.8	
		328	81.1	18.9	97.5	95.0	4.7	
30	3	0	0.0	100.0	50.0	0.0		
		16	10.9	89.1	N.A.			
		40	26.2	73.8	56.0	12.0	2.3	
		112	39.1	60.9	71.9	43.8	8.1	
		162	53.3	46.7	84.7	69.4	8.3	
		280	66.9	33.1	96.0	92.0	8.0	
		328	69.9	30.1	96.2	92.4	6.9	
35	4	0	0.0	100.0	50.0	0.0		
		16	9.8	90.2	N.A.			
		40	24.8	75.2	54.0	8.0	1.8	
		112	34.0	66.0	68.0	36.0	7.9	
		162	48.7	51.3	78.5	57.0	7.0	
		280	68.0	32.0	95.1	90.2	7.0	
		328	70.6	29.4	96.8	93.6	7.0	
Reaction system: 0.1 M SQ 32,979 (22.42 mg/ml), 0.4 M alcohol (indicated above) in 25 ml toluene; 25 uL (0.1 %v/v) deionized H ₂ O added to reaction mixture.								
Enzyme: 1.0 g (40 mg/ml) Amano lipase P-30 (from <i>Pseudomonas fluorescens</i>).								
Conditions: 28 °C, 280 rpm.								

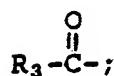
Claims

1. A process for preparing S-enantiomers of the formula

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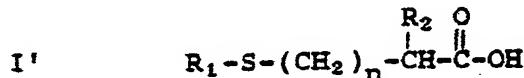


10 where R_1 is hydrogen or



15 R_2 and R_3 are each independently selected from alkyl, cycloalkyl, aralkyl or aryl; and n is 1 or 2; which process comprises reacting a racemic mixture of a compound having the formula

20



with a compound of the formula

25 VIII R_4-OH

wherein R_1 is alkyl, cycloalkyl, aryl or aralkyl, in the presence of an enzyme or microorganism capable of catalyzing the stereoselective esterification of compounds of formula I' to provide a solution containing R-enantiomeric esters and S-enantiomeric unreacted acids of formula I; and separating and recovering the S-enantiomers of formula I from said solution.

30 2. The process of claim 1 wherein R_4-OH is selected from methanol, benzyl alcohol and 1-octanol.

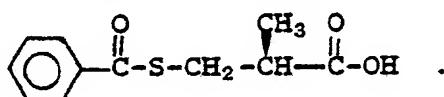
3. The process of claim 1 wherein said enzyme is selected from esterases and lipases.

4. The process of claim 1 wherein the genera of microorganisms are selected from *Mucor*, *Escherichia*, *Staphylococcus*, *Agrobacterium*, *Rhizopus*, *Aspergillus*, *Nocardia*, *Streptomyces*, *Trichoderma*, *Candida*, *Rhodotorula*, *Torulopsis*, *Bacillus*, *Alcaligenes*, *Pseudomonas*, *Brevibacterium*, *Enterobacter*, *Chromobacterium*, *Arthrobacter*, *Microbacterium*, *Mycobacterium*, *Saccharomyces*, *Penicillium*, *Botrytis*, *Chaetomium*, *Ophiobolus* and *Cladosporium*.

35 5. The process of claim 4 wherein said microorganisms are selected from *Pseudomonas fluorescens*, *Pseudomonas putida*, *Pseudomonas-ovalis*, *Escherichia coli*, *Staphylococcus aureus*, *Alcaligenes faecalis*, *Streptomyces griseus*, *Streptomyces clavuligerus*, *Nocardia erythropolis*, *Nocardia asteroides*, *Mycobacterium phlei*, *Agrobacterium radiobacter*, *Aspergillus niger* and *Rhizopus oryzae*.

40 6. The process of claim 1 wherein said enzymes are selected from Amano P (*Pseudomonas fluorescens*) which is preferred, Amano AY-30 (*Candida cylindracea*), Amano N (*Rhizopus niveus*), Amano R (*Penicillium sp.*), Amano FAP (*Rhizopus oryzae*), Amano AP-12 (*Aspergillus niger*), Amano MAP (*Mucor meihel*), Amano GC-4 (*Geotrichum candidum*), Sigma L-0382 (porcine pancreas), Sigma L-3001 (Wheat germ), Sigma L-1754 (*Candida cylindracea*), Sigma L-0763 (*Chromobacterium viscosum*) and Amano K-30 (*Aspergillus niger*). Additionally, enzymes derived from animal tissue include esterase from pig liver, α -chymotrypsin and pancreatin from pancreas.

45 7. The process of any preceding claim wherein said reaction is in the presence of an organic solvent.
8. The process of claim 7 wherein said reaction is further in the presence of an amount of water of from about 0.2 mg to about 100 mg of water per milliliter of solvent.
50 9. The process of any preceding claim wherein the product of formula I is

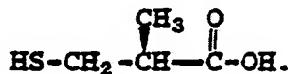


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10. The process of any one of claims 1-8 wherein the product of formula I is



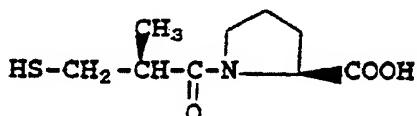
5 11. The process of any one of claims 1-8 wherein the product of formula I is



10

12. A process for the preparation of a compound of the formula

15



20 which comprises coupling a resolved carboxylic acid of the formula



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with a compound of the formula

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or its chemical equivalent,
and thereafter removing the

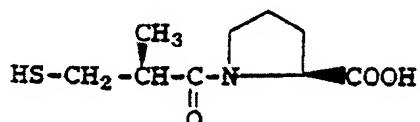
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40 moiety to provide the desired product; wherein said resolved carboxylic acid is prepared by the process of
any one of claims 1-8, R₃ being as defined in claim 1.

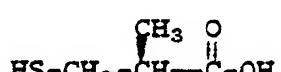
13. A process for the preparation of a compound of the formula

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which comprises coupling a resolved carboxylic acid of the formula

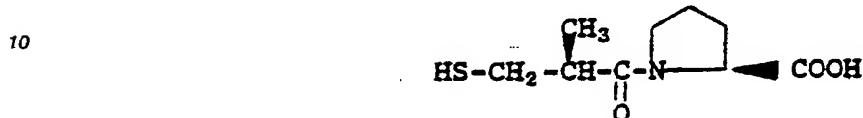


55

with a compound of the formula



- 5 or its chemical equivalent,
wherein said resolved carboxylic acid is prepared by the process of any one of claims 1-8.
14. A process for the preparation of a compound of the formula



- 15 which comprises removing the acetyl group from a resolved carboxylic acid of the formula



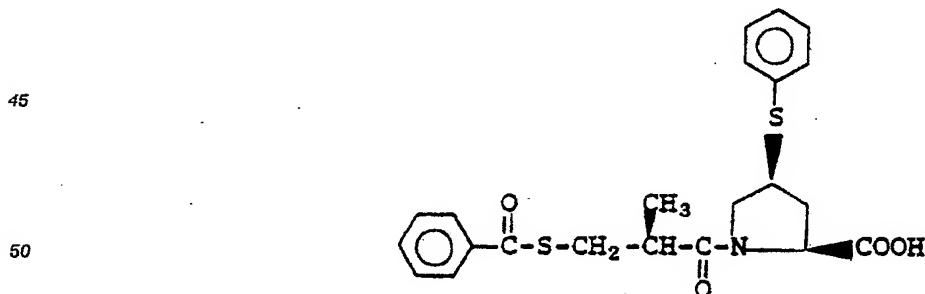
- 20 and dehydrating the so-treated acid to form a thiolactone of the formula



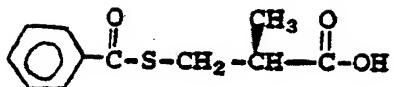
- 30 and thereafter acylating a compound of the formula.



- with said thiolactone to provide the desired product, wherein said resolved carboxylic acid is prepared by
40 the process of any one of claims 1-8, R₃ being as defined in claim 1.
15. A process for the preparation of a compound of the formula

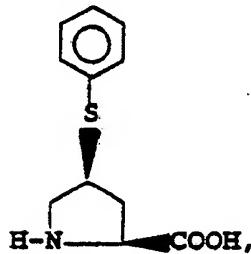


- 55 which comprises coupling a resolved carboxylic acid of the formula



5 with a compound of the formula

10

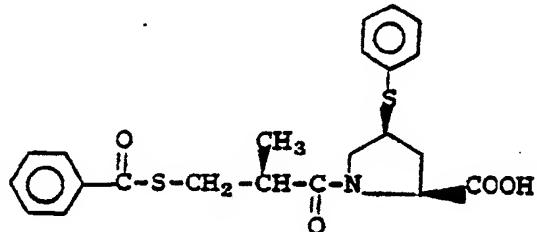


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or its chemical equivalent,
which process is further characterized in that said resolved carboxylic acid is prepared by the process of
20 any one of claims 1-8.

16. A process for the preparation of a compound of the formula

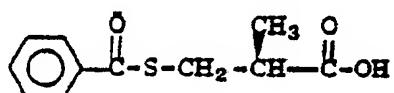
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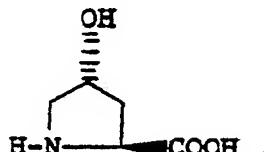
which comprises coupling a resolved carboxylic acid of the formula

35



40 with a compound of the formula

45



including esters or protected forms thereof,
50 and treating the so-formed intermediate with a suitable sulfide or disulfide to provide the desired product;
which process is further characterized in that said resolved carboxylic acid is prepared by the process of
any one of claims 1-8.

55



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number

EP 90 31 0275

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
X	PATENT ABSTRACTS OF JAPAN, vol. 13, no. 49 (C-565)[3397], 3rd February 1989; & JP-A-63 245 695 (SHOWA SHELL SEKIYU K.K.) 12-10-1988 * Abstract * ---	1-7	C 12 P 41/00 C 12 P 11/00 C 07 D 207/16
Y	WO-A-8 705 328 (WISCONSIN ALUMNI) * Pages 1-5; claims * ---	1,12,13	
Y	EP-A-0 172 614 (MITSUBISHI) * Claims * ---	1,12,13	
A	EP-A-0 130 752 (MITSUBISHI) * Claims * -----	1	
TECHNICAL FIELDS SEARCHED (Int. Cl.5)			
C 12 P C 07 D			
The present search report has been drawn up for all claims			
Place of search	Date of completion of the search	Examiner	
THE HAGUE	14-01-1991	DELANGHE L.L.M.	
CATEGORY OF CITED DOCUMENTS			
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	